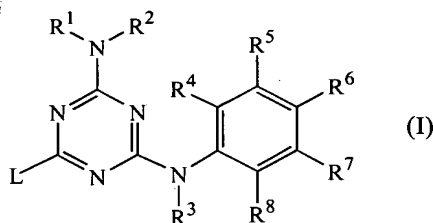


ABSTRACT

5

SUBSTITUTED DIAMINO-1,3,5-TRIAZINE DERIVATIVES

10 This invention concerns the compounds of formula



the pharmaceutically acceptable acid addition salts and the stereochemically isomeric forms thereof, wherein R¹ and R² are each independently selected from hydrogen; hydroxy; amino; optionally substituted C₁₋₆alkyl; C₁₋₆alkyloxy; C₁₋₆alkylcarbonyl; C₁₋₆alkyloxycarbonyl; Ar¹; mono- or di(C₁₋₆alkyl)amino; mono- or di(C₁₋₆alkyl)-aminocarbonyl; dihydro-2(3*H*)-furanone; or R¹ and R² taken together may form pyrrolidinyl, piperidinyl, morpholinyl, azido or mono- or di(C₁₋₆alkyl)amino-C₁₋₄alkylidene; R³ is hydrogen, Ar¹, C₁₋₆alkylcarbonyl, C₁₋₆alkyl, C₁₋₆alkyloxy-carbonyl, C₁₋₆alkyl substituted with C₁₋₆alkyloxycarbonyl; and R⁴, R⁵, R⁶, R⁷ and R⁸ are each independently selected from hydrogen, halo, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl or trihalomethyloxy; L is optionally substituted C₁₋₁₀alkyl; C₃₋₁₀alkenyl; C₃₋₁₀alkynyl; C₃₋₇cycloalkyl; Ar¹ is optionally substituted phenyl; for the manufacture of a medicine for the treatment of subjects suffering from HIV (Human Immunodeficiency Virus) infection. It further relates to new compounds being a subgroup of the compounds of formula (I), their preparation and compositions comprising them.